In the Claims:

1. (Currently Amended) A compounds having the structure of Formula I

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$$\begin{array}{c|c} H & O \\ \hline & N-A-N \\ \hline & N-R \\ \end{array}$$

FORMULA - I

its and their pharmaceutically acceptable salts, amides, enantiomers, diastereomers, N-oxides, prodrugs, metabolites or their polymorphs, wherein A is a straight or branched C_1 - C_4 alkyl chain; R is cinnamyl, benzyl, substituted benzyl, phenyl, mono- or disubstituted phenyl group substituted with the substituents independently selected from the group consisting of halogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group, or R is cinnamyl, benzyl, substituted benzyl, mono- or disubstituted phenyl group substituted with the substituents independently selected from the group consisting of halogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group, or R is cinnamyl, benzyl, substituted benzyl, mono- or disubstituted phenyl group substituted phenyl group substituted phenyl group substituted phenyl group consisting of halogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group, or R is cinnamyl, benzyl, substituted benzyl, R is cinnamyl, benzyl, substituted benzyl, mono- or disubstituted phenyl group substituted phenyl group substituted benzyl, R is cinnamyl, benzyl, substituted benzyl, R is cinnamyl, benzyl, substituted benzyl, R is cinnamyl, R is cinnamyl, benzyl, substituted benzyl, R is cinnamyl, R is cinnamyly R is cinnamyl, R is cinnamyl, R is cinnamyl, R is cinna

- 2. Cancelled
- 3. (Currently Amended) The Compounds according to claim 1 selected from the group consisting of:
 - 2-[3-{4-(2-Methoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 01);
 - 2-[3-{4-(3-Chlorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H) dione (Compound-02).
 - 2 [3 {4 (2-Methylphenyl)piperazin-1-yl}propyl] 3a,4,7,7a tetrahydro 1H-isoindole 1,3(2H) dione (Compound 03).

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- 2-[3-{4-(4-Fluorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 04);

- 2-[3-{4-(3-Trifluoromethylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 05);
- 2-[3-{4-(2-Fluorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 06);
- 2-[3-{4-(3,4-Dimethylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 07);
- 2-[3-{4-(2-Methoxy-5-fluorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 08);
- 2-[3-{4-(2-Ethylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 09);
- 2-[3-{4-(2,4-Difluorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 10);
- 2-[3-{4-(2-Ethoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 11);
- 2-[3-{4-(2-Methyl-5-chlorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 12);
- 2 [3 {4 (Phenyl)piperazin 1 yl}propyl] 3a,4,7,7a tetrahydro 1H isoindole 1,3(2H) dione (Compound 13).
- 2-[3-{4-(Benzyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 14);
- 2-[3-{4-(Cinnamyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 15);
- 2-[3-{4-(4-Nitrophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 16);
- 2-[3-{4-(3-Chloro-4-methylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 17);
- 2-[3-{4-(4-Fluoro-2-methoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 18);



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- 2-[3-{4-(Bis-4-fluorophenyl)methylpiperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 19);

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- 2-[3-{4-(2,4-Dichlorophenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 20);
- 2-[3-{4-(2,4-Dimethoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 21);
- 2-[3-{4-(2,6-Dimethylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 22);
- 2-[3-{4-(2-Isopropoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 23);
- 2-[3-{4-(2-Propoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 24);
- 2-[3-{4-(2-n-Hexyloxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 25);
- 2-[3-{4-(2,5-Dimethoxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 26);
- 2-[3-{4-(4-tert-Butylphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 27);
- 2-[3-{4-(2-Methoxy-6-hydroxyphenyl)piperazin-1-yl}propyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 28);
- 2-[3-{4-(2-Methoxyphenyl)piperazin-1-yl}-3-methylpropyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 29);
- 2-[3-{4-(2-Methoxyphenyl)piperazin-1-yl}-2-methylpropyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 30);
- 2-[3-{4-(2-Ethoxyphenyl)piperazin-1-yl}-3-methylpropyl]-3a,4,7,7a-tetrahydro-1H-isoindole-1,3(2H)-dione (Compound 31).

4. (Cancelled)

5. (Currently Amended) A method for treating benign prostatic hyperplasia in a mammal comprising administering to said mammal a compound having the structure of Formula I



$$\begin{array}{c|c} H & O \\ \hline & N-A-N \\ \hline & N-R \end{array}$$

FORMULA - I

or its pharmaceutically acceptable salts, amides, enantiomers, diastereomers, N-oxides, prodrugs, metabolites or their polymorphs, wherein A is a straight or branched C_1 - C_4 alkyl chain; R is cinnamyl, benzyl, substituted benzyl, phenyl, mono- or disubstituted with the substituents independently selected from the group consisting of halogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group, or (dihalodiphenyl) methyl.

- 6. (Currently Amended) A pharmaceutical composition comprising the <u>a</u> compound of claims 1 or 3 and a pharmaceutical acceptable carrier.
- 7. (Cancelled)
- 8. (Currently Amended) A method for treating benign prostatic hyperplasia in a mammal comprising the step of administering to the said mammal, the pharmaceutical composition according to claim 6.
- 9. (Currently Amended) A process for preparing a compounds of Formula I

$$\begin{array}{c|c}
H & O \\
N-A-N & N-R
\end{array}$$

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or its and their pharmaceutically acceptable salts, amides, enantiomers, diastereomers, N-oxides, prodrugs, metabolites or their polymorphs, wherein A is a straight or branched C_1 - C_4 alkyl chain; R is cinnamyl, benzyl, substituted benzyl, phenyl, mono- or disubstituted phenyl group substituted with the substituents independently selected from the group consisting of halogen, hydroxy, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group, or (dihalodiphenyl) methyl, which comprises comprising reacting a compound cis 1,2,3,6-tetrahydrophthalic anhydride of Formula II

FORMULA II

with piperazine derivatives 1-amino-4-substituted piperazinyl alkane of Formula III,

FORMULA III

as shown in Scheme I wherein A and R are the same as defined above, in the presence of a solvent selected from the group consisting of pyridine, n-butanol, benzene and xylene.

10. (Currently Amended) A process for preparing a compounds of Formula I

$$N-A-N$$
 $N-R$

FORMULA - I

or its and their pharmaceutically acceptable salts, amides, enantiomers, diastereomers, N-oxides, prodrugs, metabolites or their polymorphs, wherein A is a straight or branched C₁-C₄ alkyl chain;

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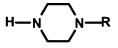
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R is cinnamyl, benzyl, substituted benzyl, phenyl, mono- or disubstituted phenyl group substituted with the substituents independently selected from the group consisting of halogen, hydroxy, C₁-C₆ alkyl, C₁-C₆ alkoxy, trifluoromethyl, nitro, trifluoroalkoxy group or(dihalodiphenyl) methyl, which comprises comprising reacting

1-(ω-haloalkyl)cis-3a,4,7,7a-tetrahydrophthalimide of Formula IV, wherein A is the same as

FORMULA IV

defined above, with a compound 1-substituted piperazine of Formula V,



FORMULA V

wherein R is the same as defined above. as shown in Scheme II.